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AMENDMENTS TO THE CLAIMS

1. (Presently amended) An aqueous solution comprising from about  $1 \times 10^{-6}$  to about  $1 \times 10^{-3}$  mol/liter of an azo dye selected from the group consisting of amaranth and Evans blue, a borate buffer and one or more masking agents, wherein the azo dye changes its coloration or coloration intensity in the presence of chlorine dioxide.
2. (Cancelled)
3. (Cancelled)
4. (Original) The aqueous solution according to claim 1, wherein the azo dye is present at a concentration of between about  $2 \times 10^{-5}$  and about  $8 \times 10^{-4}$  mol/liter.
5. (Original) The aqueous solution according to claim 1, wherein the masking agent is aqueous ammonia.
6. (Original) The aqueous solution according to claim 1, wherein the wherein the borate is present at a concentration of between about  $5 \times 10^{-3}$  and about  $1 \times 10^{-1}$  mol/liter.
7. (Original) The aqueous solution according to claim 1, further comprising one or more metal-chelating agents.
8. (Original) The aqueous solution according to claim 7, wherein the metal-chelating agent is a sodium salt of EDTA.
9. (Original) The aqueous solution according to claim 8, wherein the sodium salt of EDTA is present at a concentration of between about 0.5 and about 2 g/liter.
10. (Original) The aqueous solution according to claim 9, wherein aqueous solution contains about  $5 \times 10^{-2}$  mol/l of borate, about  $1.5 \times 10^{-2}$  mol/l of aqueous ammonia as the masking

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agent, about 1 g/l of sodium salt of EDTA and about  $2 \times 10^{-4}$  mol/l of amaranth as the azo dye.

11. (Presently presented) An aqueous solution comprising Evans blue azo dye present at a concentration of about  ~~$5 \times 10^{-4}$~~   $5 \times 10^{-5}$  mol/l, a borate buffer, aqueous ammonia present at a concentration of about  $1.5 \times 10^{-2}$  mol/l, and a sodium salt of EDTA present at a concentration of about 1 g/l, wherein the aqueous solution contains about  $5 \times 10^{-2}$  mol/l of borate and the azo dye changes its coloration or coloration intensity in the presence of chlorine dioxide.
12. (Presently amended) A process for manufacturing the aqueous solution according to one of claims 1 and 4 through 11, comprising the steps of:
  - (a) introducing the azo dye, the masking agent(s) and the borate buffer solution into a container containing a sufficient amount of double-deionized water;
  - (b) optionally adding the chelating agent predissolved in double-deionized water with stirring; and
  - (c) making up the solution to a desired volume with double-deionized water
13. (Original) The process according to claim 12 further comprising the steps of:
  - (i) dissolving the azo dye in double-deionized water;
  - (ii) introducing into a container the solution prepared in (i), followed by a borate buffer solution, and finally a solution containing one or more masking agents.
  - (iii) adding double-deionized water and measuring pH;
  - (iv) adjusting pH to about 9.2, if necessary, using concentrated aqueous ammonia solution;
  - (v) optionally adding the chelating agent with stirring; and
  - (vi) making up the solution to a desired volume with double-deionized water.

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14. (Original) The process according to claim 13, wherein the concentrated aqueous ammonia solution at about 28% (w/w) is used in the step (ii) as a masking agent and in the step (iv).
15. (Original) The process according to claim 14 further comprising the steps of:
- (i) dissolving either about 121.2 mg of amaranth [Ref. A-1016(97) Sigma] or about 56.5 mg of Evans blue (Ref. 20,633-4, Aldrich) in about 100 ml of double-deionized water;
  - (ii) dissolving about 3.09 g of boric acid in 500 ml of 0.1 M KCl solution and mixing it to homogeneity to prepare about  $5 \times 10^{-2}$  M borate buffer;
  - (iii) successively introducing into a one-liter flask, the amaranth or Evans blue solution prepared in (i), the borate buffer solution prepared in (ii), and about 1 ml of about 28% (w/w) aqueous ammonia solution;
  - (iv) adding double-deionized water and measuring pH;
  - (v) adjusting pH to about 9.2 using about 28% (w/w) aqueous ammonia solution;
  - (vi) adding about 1 g of sodium salt of EDTA with stirring until it completely dissolves; and
  - (vii) transferring the solution prepared in (vi) into a 1000-ml graduated flask and making up the total volume to the graduation mark with double deionized water.
16. (Original) A process for determining a residual chlorine dioxide content in industrial water or drinking water after treatment or in distribution circuits, comprising the steps of:
- placing the water to be analyzed in contact with the aqueous solution prepared by the process according to claim 12; and
  - measuring an absorbance of the resultant solution using a UV-visible spectrophotometer at a specific wavelength of the azo dye chosen.
17. (Original) The process according to claim 16, wherein a volume ratio:
- the water to be analyzed/the aqueous solution is between about 10 and about 30.

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18. (Presently presented) A process for determining a residual chlorine dioxide content in industrial water or drinking water after treatment or in distribution circuits, comprising the steps of:

placing the water to be analyzed in contact with the aqueous solution of claim 11 or 22, wherein a volume ratio:

the water to be analyzed / the aqueous solution is between about 10 and about 30;  
and

- measuring an absorbance of the resultant solution using a UV-visible spectrophotometer at a specific wavelength of the azo dye chosen.

19. (Presently amended) The process according to claim 18, wherein about 10 ml of the aqueous solution are placed into a 250-ml graduated flask and made up to the graduation mark with the water to be analyzed; and

an absorbance of the resultant solution is measured using a UV-visible spectrophotometer at 521 nm for amaranth or at 606 nm for Evans blue.

20. (Original) The process according to claim 19, wherein an absorbance is measured using, as a reference, the water to be analyzed to which purified and crystallized sodium thiosulphate has been added in excess of the amount required to reduce any oxidizing agents present in the water.

21. (Previously presented) The aqueous solution according to claim 1, wherein the azo dye is amaranth.

22. (New) An aqueous solution comprising amaranth azo dye present at a concentration of about  $2 \times 10^{-4}$  mol/l, a borate buffer, aqueous ammonia present at a concentration of about  $1.5 \times 10^{-2}$  mol/l, and a sodium salt of EDTA present at a concentration of about 1 g/l, wherein the aqueous solution contains about  $5 \times 10^{-2}$  mol/l of borate and the azo dye changes its coloration or coloration intensity in the presence of chlorine dioxide.